

K823052 THE 500 MONITOR SERIESNov 30, 1982
43 days to decisionK823052 · Product code: **DXN** · CardiovascularSource: <https://www.510kdatabase.net/k823052/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Oct 18, 1982
Decision date	Nov 30, 1982
Days to decision	43 days
Third-party review	No

APPLICANT

Company	Vitatek
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1981-1982

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Device record: <https://www.510kdatabase.net/k823052/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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